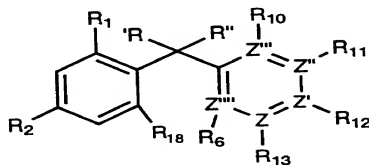
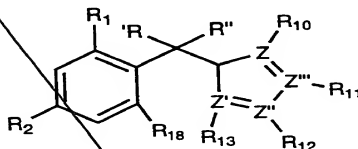


We claim:

1. A compound having the formula:



or



wherein

- 5  $R_1$  and  $R_2$ , each independently, represent hydrogen or lower alkyl or acyl having 1-4 carbon atoms;
- $R'$  and  $R''$  represent hydrogen, lower alkyl or acyl having 1-4 carbon atoms, OH, alkoxy having 1-4 carbon atoms, thiol or thio ether, or amino,
- 10 or  $R'$  or  $R''$  taken together form an oxo (keto), methano, thioketo, HO-N=, NC-N=,  $(R_7R_8)N=N$ ,  $R_{17}O-N$ ,  $R_7N$ , epoxy, cyclopropyl, or cycloalkyl group and wherein the epoxy, cyclopropyl, and cycloalkyl groups can be substituted with lower alkyl having 1-4 carbons or halogen;
- 15  $R_6$ ,  $R_{10}$ ,  $R_{11}$ ,  $R_{12}$ ,  $R_{13}$  each independently represent hydrogen, a lower alkyl having 1-4 carbons, halogen, nitro,  $OR_7$ ,  $SR_7$ ,  $NR_7R_8$  or  $(CF)_nCF_3$ , and exist only if the Z, Z', Z'', Z''', or Z''' from which it originates is C, or each independently represent hydrogen or a lower alkyl having 1-4 carbons if the Z, Z', Z'', Z''', or Z''' from

which it originates is N, and where one of R<sub>6</sub>, R<sub>10</sub>, R<sub>11</sub>, R<sub>12</sub> or R<sub>13</sub> is X;

R<sub>7</sub> represents hydrogen or a lower alkyl having 1-6 carbons;

R<sub>8</sub> represents hydrogen or a lower alkyl having 1-6 carbons;

R<sub>9</sub> represents a lower alkyl having 1-4 carbons, phenyl, aromatic alkyl, or q-hydroxyphenyl, q-bromophenyl, q-chlorophenyl, q-fluorophenyl, or q-iodophenyl, where q=2-4;

R<sub>17</sub> represents hydrogen, lower alkyl having 1-8 carbons, alkenyl (including halogen, acyl, OR, and SR, substituted alkenes),

R<sub>9</sub>, alkyl carboxylic acid (including halogen, acyl, OR, and SR, substituted alkyls), alkenyl carboxylic acid (including halogen, acyl, OR, and SR, substituted alkenes), alkyl amines (including halogen, acyl, OR, and SR, substituted alkyls), and alkenyl amines (including halogen, acyl, OR, and SR, substituted alkenes);

R<sub>18</sub> represents hydrogen, a lower alkyl having 1-4 carbons, halogen, nitro, OR, SR, NR, R<sub>8</sub>, or (CF)<sub>3</sub> CF<sub>3</sub>;

X is COOH, tetrazole, PO<sub>3</sub>H, SO<sub>3</sub>H, CHO, CH<sub>2</sub>OH, CONH<sub>2</sub>, COSH, COOR, COSR, CONHR, or COOW where W is a pharmaceutically acceptable salt, and where X can originate from any C or N on the ring;

Z, Z', Z'', Z''' and Z''', each independently, represent C, S, O, N, or a pharmaceutically acceptable salt, but is not O or S if attached by a double bond to another such Z or if attached to another such Z which is O or S, and is not N if attached by a single bond to another such Z which is N, and

n = 0-3.

2. A compound of claim 1 wherein said compound selectively activates Retinoid X Receptors in preference to Retinoic Acid Receptors.

3. A compound selected from the group consisting of 4-  
[1-(2-methyl-4-t-butylphenyl)ethenyl] benzoic acid,  
4-[1-(2-methyl-4-t-butylphenyl)cyclopropyl] benzoic acid,  
4-[(2-methyl-4-t-butylphenyl)carbonyl] benzoic acid,  
5 4-[(2-methyl-4-t-butylphenyl)carbonyl] benzoic acid oxime, and  
4-[1-(2-methyl-4-t-butylphenyl)carbonyl] benzoic acid

~~methyloxime~~

4. A pharmaceutical composition comprising in a  
pharmaceutically acceptable vehicle suitable for enteral,  
10 parenteral, or topical administration, one or more compound of  
claim 1.

5. A method for modulating a process mediated by one or  
more Retinoid X Receptors, said method comprising causing said  
process to be conducted in the presence of at least one compound as  
15 ~~set forth in claim 1.~~

6. A method according to claim 5 wherein said process  
is the *in vivo* modulation of lipid metabolism, *in vivo* modulation of  
skin-related processes, *in vivo* modulation of malignant cell  
development, *in vivo* modulation of premalignant lesions, or *in vivo*  
20 modulation of programmed cell death.

7. A method according to claim 5 wherein said process  
is in *in vivo* or *in vitro* cellular growth and differentiation, or  
*in vivo* limb morphogenesis.

8. A method for modulating a process mediated by one or  
25 more Retinoid X Receptors, said method comprising administering to

a mammalian subject an amount, effective to modulate said process mediated by said one or more Retinoid X Receptors, of one or more compound of claim 1.

9. A method for treating a mammalian subject requiring Retinoid X Receptor therapy comprising administering to such subject a pharmaceutically effective amount of one or more compounds as set forth in claim 1.

10. A method for increasing plasma concentrations of high density lipoprotein in a mammalian subject comprising administering to such subject a pharmaceutically effective amount of one or more compounds as set forth in claim 1.

11. A method for modulating a process mediated by intracellular receptors, said method comprising causing said process to be conducted in the presence of a composition comprising a first compound as set forth in claim 1 which selectively activates Retinoid X Receptors in preference to Retinoid Acid Receptors, in combination with a second compound which activates one or more intracellular receptors other than Retinoid X Receptors, and wherein the physiological effect in mammals produced by said composition at a given concentration is greater than the additive effect achieved by utilizing each said compound alone at said concentration.

add  
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